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*1 / 1 PLUSPAT - ©QUESTEL-ORBIT***Patent Number :** US6063806 A 20000516 [US6063806]**Title :**

(A) Indolyl or indolinyl derivatives and medicinal use thereof as ACAT or lipid peroxidation inhibitors

Patent Assignee :

(A) KYOTO PHARMA IND (JP)

Patent Assignee :

Kyoto Pharmaceutical Industries, Ltd., [JP]

Inventor(s) :

(A) KAMIYA SHOJI (JP); SHIRAHASE HIROAKI (JP); MATSUI HIROSHI (JP); NAKAMURA SHOHEI (JP); WADA KATSUO (JP)

Application Nbr :

US5120298 19980403 [1998US-0051202]

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Intl Patent Class :

(A) A61K-031/40 C07D-209/08 C07D-209/12 C07D-209/14 C07D-209/18

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K. Yee et al., "Novel Series of Selective Leukotriene Antagonists: Exploration and Timization of the Acidic Region in 1,6-Disubstituted Indoles and Indazoles", Journal of

Medicinal Chemistry, vol. 33, No. 9, pp. 2437-2451, 1990.

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F. Brown et al., "Evolution of a Series of Peptidoleukotriene Antagonists: Synthesis and Structure-Activity Relationships of 1,6-Disubstituted Indoles and Indazoles", Journal of Medicinal Chemistry, vol. 33, No. 6, pp. 1771-1781, 1990.

Publication Stage :

(A) United States patent

Abstract :


A heterocyclic derivative of the formula (I) (Chemical Structure image 'I' not included in text) wherein each symbol is as defined in the specification, and pharmaceutically acceptable salts thereof. The compound (I) of the present invention and pharmaceutically acceptable salts thereof exhibit superior ACAT inhibitory activity and lipoperoxidation inhibitory activity in mammals, and are useful as ACAT inhibitors and lipoperoxidation inhibitors. Specifically, they are useful for the prophylaxis and treatment of arteriosclerosis, hyperlipemia, arteriosclerosis in diabetes, and cerebrovascular and cardiovascular ischemic diseases.

Update Code :

2000-22

1 / 1 LGST - ©EPO

Patent Number :

 US6063806 A 20000516 [US6063806]

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US5120298 19980403 [1998US-0051202]

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2003-46

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051202 (09) 6063806 May 16, 2000

UNITED STATES PATENT AND TRADEMARK OFFICE GRANTED PATENT

6063806

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May 16, 2000

Indolyl or indolynyl derivatives and medicinal use thereof as ACAT or lipid peroxidation inhibitors

REISSUE: June 27, 2003 - Reissue Application filed Ex. Gp.: 1614; Re. S.N. 10/609,224 (O.G. November 4, 2003)

APPL-NO: 051202 (09)

FILED-DATE: April 3, 1998

GRANTED-DATE: May 16, 2000

CORE TERMS: sub, sup, compound, br-t, indoline, evaporated, mixture, alkyl, dissolved, residue ...

ENGLISH-ABST:

A heterocyclic derivative of the formula (I) ##STR1## wherein each symbol is as defined in the specification, and pharmaceutically acceptable salts thereof. The compound (I) of the present invention and pharmaceutically acceptable salts thereof exhibit superior ACAT inhibitory activity and lipoperoxidation inhibitory activity in mammals, and are useful as ACAT inhibitors and lipoperoxidation inhibitors. Specifically, they are useful for the prophylaxis and treatment of arteriosclerosis, hyperlipemia, arteriosclerosis in diabetes, and cerebrovascular and cardiovascular ischemic diseases.

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Date/Time: Wednesday, May 12, 2004 - 4:10 PM EDT


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